AMENDMENTS TO THE CLAIMS:

Please amend claim 68, as shown below.

This listing of claims will replace all prior versions and listings of claims in the Application:

Claims 1-35 (cancelled)

Claim 36 (previously presented): A compound having the formula:

Ia

wherein:

R₁ represents a radical selected from the group consisting of

$$\begin{array}{c} \stackrel{R_2}{\longleftarrow} \stackrel{R_2}{\longleftarrow} \stackrel{R_3}{\longleftarrow} R_4 \\ \stackrel{R_3}{\longleftarrow} , -NR_5R_6, -SO_2NR_7R_8, \ \text{hydroxyalkyl}, \ \text{hydroxyalkoxy, polyfluoroalkyl}, \ \text{dialkylaminoalkyl}, \ R_9, -OR_9, \end{array}$$

$$-N$$
 $SO_2^-R_{10}$
 $SO_2^-R_{11}$; n being an integer from 1 to 4;

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R₂ and R₃ are each independently selected from the group consisting of straight or branched chain alkyl and hydrogen;

 R_4 is a radical selected from the group consisting of a substituted or unsubstituted phenyl radical, an unsubstituted or substituted heterocyclic radical, and $-NR_{12}R_{13}$;

R₅ and R₇ are independently selected from the group consisting of alkoxyalkyl, hydroxyalkyl, polyhydroxyalkyl, aralkyl, R₉, -(C=O)R₁₄ and -(C=O)R₉;

 R_6 , R_8 , R_{12} , and R_{13} are independently selected from the group consisting of hydrogen, alkyl, alkoxyalkyl, polyfluoroalkyl, hydroxyalkyl, polyhydroxyalkyl, aralkyl, R_9 , -(C=O) R_{15} and -(C=O) R_9 ;

or R₅ and R₆ taken together with the nitrogen to which they are attached form a substituted or unsubstituted heterocyclic radical, said heterocyclic radical optionally containing one to two additional heteroatoms independently selected from the group consisting nitrogen, oxygen, and sulfur;

or R₇ and R₈ taken together with the nitrogen to which they are attached form a substituted or unsubstituted heterocyclic radical, said heterocyclic radical optionally containing one to two additional heteroatoms independently selected from the group consisting nitrogen, oxygen, and sulfur;

said phenyl and heterocyclic radical substituents being at least one selected from the group consisting of alkyl, amino, hydroxy, carbonyl, monoalkylamino, dialkylamino, halogen, and alkoxy;

R₉ is a radical of the formula –W-O(C=O)-CH₃, W being a straight- or branched- chain alkylene group of 1 to 6 carbon atoms;

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 R_{10} and R_{11} are radicals independently selected from the group consisting of alkyl, halo, haloalkyl, and polyfluoroalkyl;

R₁₄ is a hydroxyalkyl, alkoxyalkyl or cycloalkyl group;

 R_{15} is an alkyl, hydroxyalkyl, alkoxyalkyl or cycloalkyl group, and pharmaceutically acceptable salts of said compound.

Claim 37 (previously presented): A compound according to claim 36, selected from the group consisting of:

- 2,2'-[[3-(2,2,2-Trifluoroethyl)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-(4-Morpholinyl)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-(1-Piperidinyl)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-(4-Morpholinylsulfonyl)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-(Methoxyethoxy)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[[3-Bis(phenylmethyl)amino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[1-(4-Methylpiperazinyl)methyl]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-(Diethylaminomethyl)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;

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- 2,2'-[[3-(Dimethylaminomethyl)phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[4-(Morpholinyl)methyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(4-Hydroxybutyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[2-(1,1-Dioxide-2,3,4,5-tetrahydro)isothiazolyl]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(3-Hydroxypropyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(2-Hydroxyethyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[1-(4-Hydroxypiperidinyl)]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(3-Hydroxypropyl)-N-methylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(4-Acetoxybutyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(2-Hydroxyethyl)-N-methylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;
- 2,2'-[[3-[N-(4-Hydroxybutyl)-N-methylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;

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2,2'-[[3-[N-(2-Hydroxyethyl)-N-propyllamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;

2,2'-[[3-[N-(4-Hydroxybutyl)-N-propylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol;

2,2'-[[3-[N-(6-Hydroxyhexyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol; and

2,2'-[[3-[N-(5-Hydroxypentyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1-yl)imino]methyl]]phenol.

Claim 38 (previously presented): The compound according to claim 36 having the formula:

Claim 39 (previously presented): The compound according to claim 36 having the formula:

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Claim 40 (previously presented): The compound according to claim 36 having the name 2,2'-[[3-[N-(4-Hydroxybutyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1yl)imino]methyl]]phenol.

Claim 41 (previously presented): The compound according to claim 36 having the name 2,2'-[[3-[N-(2-Hydroxyethyl)-N-methylamino]phenyl]methylene]bis[4-[[(5-methyl-1Htetrazol-1-yl)imino]methyl]]phenol.

Claim 42 (previously presented): The compound according to claim 36 having the name 2,2'-[[3-[N-(2-Hydroxyethyl)-N-ethylamino]phenyl]methylene]bis[4-[[(5-methyl-1H-tetrazol-1yl)imino]methyl]]phenol.

Claim 43 (previously presented): A pharmaceutical composition for treating or preventing pneumovirus infection, said composition comprising a compound according to claim 36 in an amount effective to attenuate infectivity of said virus, and a pharmaceutically acceptable carrier medium.

Claim 44 (previously presented): A pharmaceutical composition according to claim 43, further comprising at least one supplemental active agent selected from the group consisting of

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interferon, ribavirin and an immunomodulator, an immunoglobulin, an anti-flammatory agent, an antibiotic, an anti-viral and an anti-infective.

Claim 45 (previously presented): A pharmaceutical composition according to claim 43, wherein said pharmaceutically acceptable carrier medium comprises ethanol.

Claim 46 (previously presented): A pharmaceutical composition according to claim 43, wherein said pharmaceutically acceptable carrier medium comprises propylene glycol.

Claim 47 (previously presented): A pharmaceutical composition according to claim 43, wherein said pharmaceutically acceptable carrier medium comprises water.

Claim 48 (previously presented): A pharmaceutical composition according to claim 45, wherein said pharmaceutically composition comprises at least 50% ethanol.

Claim 49 (previously presented): A pharmaceutical composition according to claim 45, wherein said pharmaceutically composition comprises at least 60% ethanol.

Claim 50 (previously presented): A pharmaceutical composition according to claim 45, wherein said pharmaceutically composition comprises at least 70% ethanol.

Claim 51 (previously presented): A pharmaceutical composition according to claim 45, wherein said pharmaceutically composition comprises at least 80% ethanol.

Claim 52 (previously presented): A pharmaceutical composition according to claim 45, wherein said pharmaceutically composition comprises at least 90% ethanol.

Claim 53 (previously presented): A pharmaceutical composition according to claim 52, wherein said pharmaceutically composition comprises less than 5% water.

Claim 54 (previously presented): A pharmaceutical composition according to claim 45, wherein said pharmaceutically acceptable carrier medium comprises ethanol, water, and propylene glycol.

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Claim 55 (previously presented): A pharmaceutical composition according to claim 54, wherein said pharmaceutically composition comprises about 85% ethanol, about 10% propylene glycol, and about 5% water.

Claim 56 (previously presented): A method of treatment of pneumovirus infection in a patient in need of said treatment, said method comprising administering to said patient a therapeutically effective amount of a compound according to claim 36.

Claim 57 (previously presented): A method as claimed in claim 56, wherein said compound is administered through inhalation or intribation.

Claim 58 (previously presented): A method as claimed in claim 56, wherein said compound is administered by an electrohydrodynamic delivery device.

Claim 59 (previously presented): A method as claimed in claim 56, wherein said electrohydrodynamic delivery device is hand-held.

Claim 60 (previously presented): A method as claimed in claim 58, wherein said electrohydrodynamic delivery device is disposable.

Claim 61 (previously presented): A method as claimed in claim 58, wherein said electrohydrodynamic delivery device is for a single user.

Claim 62 (previously presented): A method as claimed in claim 58, wherein said electrohydrodynamic delivery device comprises a removable mouthpiece.

Claim 63 (previously presented): A method as claimed in claim 58, wherein said electrohydrodynamic delivery device comprises a mask.

Claim 64 (previously presented): A method of treating cells in culture that are susceptible to infection by, or infected or contaminated with a pneumovirus, said method comprising administering to said cultures an effective amount of a compound according to claim 36.

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Claim 65 (previously presented): A method of treating biological materials that are susceptible to infection by, or infected or contaminated with a pneumovirus, said method comprising administering to said materials an effective amount of a compound according to claim 36.

Claim 66 (previously presented): A compound having the formula:

wherein:

 $R_{1}{}^{\prime}$ represents a radical selected from the group consisting of

$$\begin{array}{c} -\left(\stackrel{R_2}{\circ}\right)_{n}^{R_2} \\ +\stackrel{R_3}{\circ} \\ +NR_5R_6, -SO_2NR_7R_8, \ hydroxyalkyl, \ hydroxyalkoxy, \ polyhydroxyalkyl, \\ \\ alkoxyalkoxy, \ polyfluoroalkyl, \ dialkylaminoalkyl, \ R_9, -OR_9, \end{array}$$

$$SO_2$$
-R
 SO_2 -R
 11 ; n being an integer from 1 to 4;

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 R_2 and R_3 are each independently selected from the group consisting of straight or branched chain alkyl and hydrogen;

 R_4 is a radical selected from the group consisting of a substituted or unsubstituted phenyl radical, an unsubstituted or substituted heterocyclic radical, and $-NR_{12}R_{13}$;

 R_5 and R_7 are independently selected from the group consisting of alkoxyalkyl, hydroxyalkyl, polyhydroxyalkyl, aralkyl, R_9 , -(C=O) R_{14} and -(C=O) R_9 ;

 R_6 , R_8 , R_{12} , and R_{13} are independently selected from the group consisting of hydrogen, alkyl, alkoxyalkyl, polyfluoroalkyl, hydroxyalkyl, polyhydroxyalkyl, aralkyl, R_9 , -(C=O) R_{15} and -(C=O) R_9 ;

or R₅ and R₆ taken together with the nitrogen to which they are attached form a substituted or unsubstituted heterocyclic radical, said heterocyclic radical optionally containing one to two additional heteroatoms independently selected from the group consisting nitrogen, oxygen, and sulfur;

or R₇ and R₈ taken together with the nitrogen to which they are attached form a substituted or unsubstituted heterocyclic radical, said heterocyclic radical optionally containing one to two additional heteroatoms independently selected from the group consisting nitrogen, oxygen, and sulfur;

said phenyl and heterocyclic radical substituents being at least one selected from the group consisting of alkyl, amino, hydroxy, carbonyl, monoalkylamino, dialkylamino, halogen, and alkoxy;

R₉ is a radical of the formula –W-O(C=O)-CH₃, W being a straight- or branched- chain alkylene group of 1 to 6 carbon atoms;

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 R_{10} and R_{11} are radicals independently selected from the group consisting of alkyl, halo, haloalkyl, and polyfluoroalkyl;

R₁₄ is a hydroxyalkyl, alkoxyalkyl or cycloalkyl group;

 R_{15} is an alkyl, hydroxyalkyl, alkoxyalkyl or cycloalkyl group, and pharmaceutically acceptable salts of said compound.

Claim 67 (cancelled).

Claim 68 (currently amended): A compound having the formula:

CHO
OH
OR
$$_{b}$$
 $_{CR_{b}}$

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Ш

wherein R_b is selected from the group consisting of -CH₂OCH₃, -CH₂OCH₂CH₃, -

CH(CH₃) OCH₂CH₃, -CH₂-OCH₂CH₂-OCH₃, -CH₂-OCH₂CH₂-Si(CH₃)₃, -CH₃, - $CH_2C_6H_5$, $-(CH_2)_2Si(CH_3)_3$, $-CON(R_cR_d)_2$, $-CSN(R_cR_d)_2$, and $-PO(NR_cR_d)_2$;

R_c and R_d are independently selected from an alkyl group;

R₁" represents a radical selected from the group consisting of

$$-\left(\stackrel{\stackrel{1}{c}}{\stackrel{2}{-}}_{n}R_{4}\right)$$

$$\stackrel{R_{3}}{, -NR_{5}R_{6}, -SO_{2}NR_{7}R_{8}, \text{ hydroxyalkyl, hydroxyalkoxy, polyhydroxyalkyl,}}{\text{alkoxyalkoxy, polyfluoroalkyl, dialkylaminoalkyl, }R_{9}, -OR_{9}}$$

$$SO_2 = R_{10}$$

 $SO_2 = R_{11}$; n being an integer from 1 to 4;

R₂ and R₃ are each independently selected from the group consisting of straight or branched chain alkyl and hydrogen;

R₄ is a radical selected from the group consisting of a substituted or unsubstituted phenyl radical, an unsubstituted or substituted heterocyclic radical, and -NR₁₂R₁₃;

 R_5 and R_7 are independently selected from the group consisting of alkoxyalkyl, hydroxyalkyl, polyhydroxyalkyl, aralkyl, R₉, -(C=O)R₁₄ and -(C=O)R₉;

R₆, R₈, R₁₂, and R₁₃ are independently selected from the group consisting of hydrogen, alkyl, alkoxyalkyl, polyfluoroalkyl, hydroxyalkyl, polyhydroxyalkyl, aralkyl, R₉, -(C=O)R₁₅ and $-(C=O)R_9$;

or R₅ and R₆ taken together with the nitrogen to which they are attached form a substituted or unsubstituted heterocyclic radical, said heterocyclic radical optionally containing

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one to two additional heteroatoms independently selected from the group consisting nitrogen, oxygen, and sulfur;

or R_7 and R_8 taken together with the nitrogen to which they are attached form a substituted or unsubstituted heterocyclic radical, said heterocyclic radical optionally containing one to two additional heteroatoms independently selected from the group consisting nitrogen, oxygen, and sulfur;

said phenyl and heterocyclic radical substituents being at least one selected from the group consisting of alkyl, amino, hydroxy, carbonyl, monoalkylamino, dialkylamino, halogen, and alkoxy;

R₉ is a radical of the formula –W-O(C=O)-CH₃, W being a straight- or branched- chain alkylene group of 1 to 6 carbon atoms;

 R_{10} and R_{11} are radicals independently selected from the group consisting of alkyl, halo, haloalkyl, and polyfluoroalkyl;

R₁₄ is a hydroxyalkyl, alkoxyalkyl or cycloalkyl group; and

R₁₅ is an alkyl, hydroxyalkyl, alkoxyalkyl or cycloalkyl group.

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